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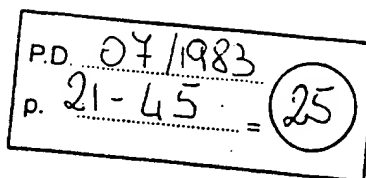
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XP-002105956

Mol. Biol. Med. (1983) 1, 21-45



**Sequence Analysis of the 17,166 Base-pair *Eco*RI fragment C of
B95-8 Epstein-Barr Virus**

A. T. Bankier, P. L. Deininger, P. J. Farrell and B. G. Barrell†

*MRC Laboratory of Molecular Biology, MRC Centre
Hills Road, Cambridge CB2 2QH, England*

(Received 4 March 1983)

Summary In order to provide a framework for understanding the molecular biology of Epstein-Barr virus (EBV), we are determining the DNA sequence of the virus and studying the organization of genes on the viral genome. In this paper we report the DNA sequence of the *Eco*RI C fragment of the B95-8 strain of EBV. The large (approximately 13.6 kb†) deletion in this strain has been located by comparison with the DNA sequence of EBV isolated from Raji cells. The sequence has been analysed for possible protein coding regions and transcriptional control sites. At least eight large open reading frames are found, some of them associated with canonical promoter and polyadenylation sequences. The sequences of some of the encoded proteins suggest that they are membrane proteins. It is known that antibodies to major membrane glycoproteins of EBV can neutralize infection in tissue culture. A possible relationship between some of the encoded proteins and the major membrane glycoproteins of the virus is discussed.

Introduction

Relatively little is known about the molecular biology of Epstein-Barr virus (EBV) despite its medical importance as the causative agent of infectious mononucleosis and its association with Burkitt's lymphoma and nasopharyngeal carcinoma (reviewed by Epstein & Achong, 1979; Tooze, 1980). An understanding of the structure of the viral genes and the control of their expression will be extremely useful in developing vaccines or synthetic immunogens against EBV infection. Since EBV transforms some human cells in tissue culture an analysis of the structure and function of the genes required to effect this transformation should make an important contribution to our understanding of natural human carcinogenesis. The virus also exhibits long-term latent infections in humans and provides a system in which to study the importance of this type of infection. The alternative growth patterns of the virus (latent and

† Author to whom all correspondence should be addressed.

‡ Abbreviation used: kb, 10³ bases.

productive infections) and their tissue specificity within the human body provide an interesting problem in the control of gene expression since different groups of viral genes are thought to be expressed in the different systems.

Progress in understanding the organization of EBV gene expression has been complicated by both the large size of the viral genome and the lack of a simple permissive cell culture system for lytic growth of the virus. The use of diterpene ester tumour promoters to induce EBV synthesis in transformed cell lines (zur Hausen *et al.*, 1978, 1979) has made it much easier to study viral gene expression and the cloning of the viral DNA has made analysis of the viral genome organization possible. Restriction enzyme maps for several enzymes (*EcoRI*, *BamHI*, *SaII* and *HindIII*) have been established and clone libraries have been made by recombinant DNA techniques for the *EcoRI* and *BamHI* restriction enzyme fragments (Dambaugh *et al.*, 1980; Skare & Strominger, 1980; Arrand *et al.*, 1981).

The major features of the genome, which is approximately 170 kbp in length, are two unique regions, the small unique region (U_S) and the large unique region (U_L), which are separated by a variable (up to 12 copies) number of direct repeats each 3.07 kb in length. Another class of direct repeats each about 540 bases in length, again varying in number, are found at both ends of the linear viral DNA. These ends can circularize to form an intracellular episomal form. An approximate transcription map has been constructed by hybridization of mRNA to the *EcoRI* and *BamHI* restriction enzyme fragments (Hummel & Kieff, 1982a). Using hybrid selected translation, the genes for some EBV proteins have been assigned to particular restriction fragments (Hummel & Kieff, 1982b). In addition the genes for two small RNAs transcribed by RNA polymerase III have been located by hybridization and sequence analysis in the small unique region (U_S). The restriction enzyme mapping and hybridization studies have revealed the existence of deletions in different strains of EBV (Pritchett *et al.*, 1975; Sugden *et al.*, 1976; Hayward & Kieff, 1977). The B95-8 cell line was established by infecting marmoset lymphocytes with EBV derived from peripheral blood leukocytes of a patient with infectious mononucleosis (Miller *et al.*, 1972). B95-8 DNA has been shown to be missing approximately 13.6 kb from near the right-hand end of the long unique region (Deliuss & Bornkamm, 1978; Raab-Traub *et al.*, 1978, 1980; Bornkamm *et al.*, 1980; Heller *et al.*, 1981). Part of this deleted region has been shown to be homologous to a region just to the right of the large internal repeats (Raab-Traub *et al.*, 1980). The significance of the deleted DNA is unknown since, so far as we know, the B95-8 strain shows all the functional properties of other strains not carrying the deletion. However, two components of the EBV membrane antigen complex (gp350 and gp220), which are antigenically related, are expressed in a uniquely abnormal ratio in the B95-8 line (Thorley-Lawson & Geilinger, 1980).

We are in the process of determining the complete DNA sequence of EBV; here we describe the DNA sequence analysis of the *EcoRI* C fragment of B95-8 DNA (see restriction map in Fig. 1) and the identification of the deletion point by comparison with sequences determined in Raji EBV DNA that does not carry the deletion. We present a detailed analysis of the possible coding regions and potential transcription sequences found in this region of EBV.

† See footnote to p. 21.

Materials and methods

(1) DNA recombinants

Recombinant cosmids containing the *EcoRI* C fragment of B95-8 and Raji EBV DNA were obtained from B. E. Griffin and J. Arrand, ICRF Laboratories, Lincoln's Inn Fields, London. DNA purifications were carried out as previously described (Birnboim & Doly, 1979). Recombinant DNA inserts were isolated by restriction enzyme digestion followed by electrophoresis through LGT agarose (Wieslander, 1979). A random subclone library of the *EcoRI* C fragment was generated by sonication (Deininger, 1983) followed by cloning into the bacteriophage M13 mp7 (Messing *et al.*, 1981), M13 mp8 or mp9 (Messing & Vieira, 1982) vector systems.

(2) DNA sequence analysis

Single-stranded DNA templates prepared from the M13 subclone library (Sanger *et al.*, 1980) were sequenced by the dideoxynucleotide chain termination method (Sanger *et al.*, 1977; Sanger & Coulson, 1978) using a complementary synthetic oligonucleotide primer (Duckworth *et al.*, 1981). Sequence data from the M13 subclone library were aligned and overlapped by computer (Staden, 1982). The complete sequence of both strands of DNA was established. Completion of the sequence required some non-random sequencing. Initially, M13 clones were not obtained surrounding an *Eco* K site, which happens to occur in the *EcoRI* C sequence (*Escherichia coli* JM101 is r_2). Clones covering this region were generated by isolation of restriction fragments covering the site and forced cloning into M13. A total of 536 sequencing gel readings were performed to obtain the *EcoRI* C sequence. The total length of sequence read was 112,581 bases. On average each nucleotide of the sequence was determined 6.5 times.

(3) Nomenclature

Because the large open reading frames and potential transcription signals are found scattered on both strands of the DNA we have used the nomenclature R (rightward) strand and L (leftward) strand. Reading frames on the R strand are named RFI-5 and those on the L strand are named LFI-6. In order to avoid confusion, in the future we propose to prefix these names with the initial letter of the restriction enzyme and the fragment number, e.g. EC - for *EcoRI* fragment C; BX - for *BamHI* fragment X. The *in vitro* promoters are designated R or L (for rightward and leftward) followed by a number. In order to avoid the confusion of using two nucleotide numbering systems for the R and L strands, we have used the R strand numbering for both strands throughout. Although most groups use a map of the type shown in Figure 1, we note

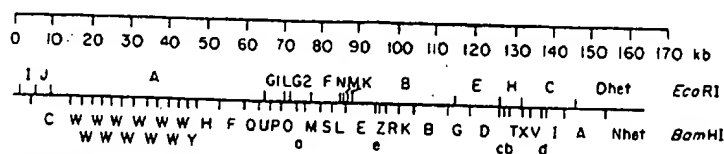


Figure 1. Restriction map of *EcoRI* and *BamHI* sites in B95-8 EBV (Arrand *et al.* 1981). Scale is in kb.

that a few authors prefer an opposite sense map of the EBV genome (Hayward *et al.*, 1980).

Results and discussion

(1) Sequence analysis

Because of the lack of genetic data for EBV, we have adopted the approach of first establishing its DNA sequence then using this in a predictive fashion to design further experiments to understand the coding and expression of the EBV genome.

As a first stage in the detailed analysis we have determined the DNA sequence of the *EcoRI* C fragment (Fig. 2). Marked on the sequence are major open reading frames, potential promoters and splice junctions, potential poly(A) addition sites (AATAAA) and the three promoters which were detected by *in vitro* transcription (Farrell *et al.*, 1983). The organization of the reading frames, promoters and poly(A) addition sites is shown in a simplified form in Figure 3. Although smaller coding regions, whether genes or individual exons, are possible we have excluded those open reading frames lacking a possible initiation codon and under 200 codons in length. We have also examined the codon usage of the reading frames to see whether they have a distinctive preference in order to try to predict coding and non-coding regions.

It is difficult to predict the function of a DNA sequence without any other biological information about it. This is especially true of eukaryotic sequences where complex RNA splicing may occur (for example, adenovirus, reviewed by Tooze, 1980). Although there are specific sequences associated with transcription and RNA splicing, these are consensus sequences and any particular signal sequence can be quite different from the canonical sequences such as the TATA box (Corden *et al.*, 1980) and the CCAAT box (Efstratiadis *et al.*, 1980) of eukaryotic promoters. In some cases nothing resembling these sequences can be found in eukaryotic promoters, e.g. SV40 and polyoma late promoters and the adenovirus E11A and IVA2 promoters (see review by Shenk, 1981). There are consensus sequences for splice donor and acceptor junctions of λ AG/GTPuAGT and Py₁N Py AG/G, respectively (see collection of sequences by Mount, 1982). The redundancy found in these sequences makes their prediction very difficult. Nevertheless, when these promoter and splice sequences are near to their consensus sequences and are present in logical combinations with open reading frames and the polyadenylation sequence AATAAA, then their presence can be highly suggestive of transcriptional units and protein coding regions.

Figure 2. The sequence of the *EcoRI* fragment C of EBV from B95-8 cells is shown in double-stranded form and numbered as for the R strand. Large open reading frames (greater than 200 codons) are translated into the one-letter amino acid code and identified in the right-hand margin. The three promoters identified by Farrell *et al.* (1983) are shown with the canonical promoter sequence TATAAAA indicated by *.....*. Other possible promoter sequences are similarly marked but are otherwise unlabelled. Possible polyadenylation sequences (AATAAA) are marked by + + + + +. Potential splice sequences that are noted or discussed in the text are shown by --X-- to denote the canonical donor sequence λ AG/GTPuAGT and by -----|| for the canonical acceptor sequence Py₁AG/G (Mount, 1982). A semi-repetitive sequence is shown by dashed overlining. Twofold symmetric sequences mentioned in the text are shown by < < < < > > > > where < and > are complementary bases. The B95-8 deletion point as compared with Raji DNA is also shown.

25

FIG. 2

27

FIG. 2, continued

[illegible]

FIG. 2. continued

29

FIG. 2, continued

FIG. 2, continued

[illegible]

31

2

FIG. 2, continued

FIG. 2. continued

33

FIG. 2, continued

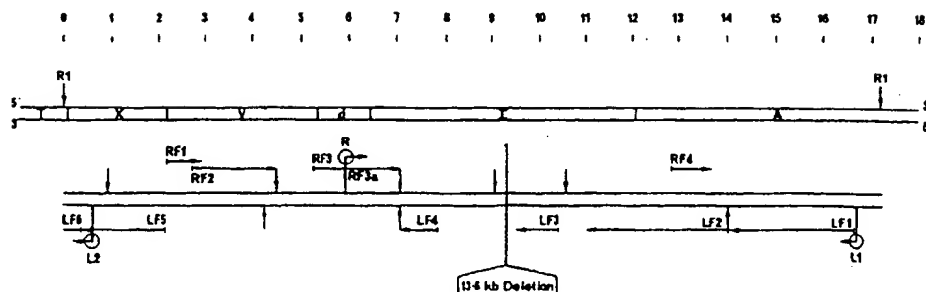


Figure 3. Organization of the *EcoRI* C fragment of B95-8 EBV. Open reading frames (RF1 etc) are shown by horizontal thin arrows, AAUAAA sequences (vertical thin arrows) and the position of the deletion relative to EBV from Raji cells are also marked. The three promoters L1, L2 and R are indicated. The scale is in kb.

(2) Codon preferences

EBV has a G + C content of 58% (Weinberg & Becker, 1969; Schultze-Holthausen & zur Hausen, 1970) and the nucleotide composition of the *EcoRI* C fragment matches this closely (A 20.6%, G 29.1%, C 29.4%, T 20.9%). The pattern of codon usage in open reading frames detected in the DNA sequence can be used to predict whether these are actually translated. For any particular amino acid, the first two positions in its codons are nearly always fixed (except for the 6 codon families: leucine UUR, CUX; serine UCX, AGY and arginine CGX and AGR), so with a fairly random amino acid sequence the composition of the nucleotides in the first two positions will also be fairly random. In a coding region with a strong G + C bias the excess G + C nucleotides will tend to be concentrated into the third position of the codons. So the nucleotide bias tends to be exaggerated in the codon usage and may produce a highly characteristic codon usage pattern. If the region is not being expressed into protein there is no such constraint pushing the excess G + C nucleotides into the third position and the codon usage pattern will be quite different. Such a highly biased codon usage pattern has been observed previously in bacteriophage ϕ X with 31% T. In a random protein sequence encoded by a nucleotide sequence with a 58% G + C content, the G + C content of the first two codon positions should be 50% and the G + C content of the third position would be predicted to be 74%. Table 1 lists the % G + C content in the third position of the codons in all the reading frames shown in Figures 2 and 3. The average G + C content in the third position of the codons in these readings frames is 69%, approaching the third position G + C content calculated for a random protein sequence and substantially higher than the 58% of the whole sequence. This highly non-random distribution of the G + C content in the reading frames make it extremely likely that they or large elements of them correspond to protein coding regions. The biased codon usage will be detected by the FRAMESCAN program (Staden & McLachlan, 1982), which analyses the codon usage of a known gene or potential gene and uses the characteristic codon pattern to predict other similar possible coding regions. Thus we can quickly scan EBV sequences for similar putative coding regions

TABLE 1
Third position G + C in reading frames

Frame	Length	3rd G + C	% 3rd G + C
EC-RF1*	186	118	63.4
EC-RF2	571	445	77.9
EC-RF3	606	425	70.1
EC-RF3a	346	250	72.5
EC-RF4†	290	173	59.6
EC-LF1	858	651	75.9
EC-LF2	1016	832	81.9
EC-LF3	313	229	73.2
EC-LF4	249	144	57.8
EC-LF5	608	369	60.7
EC-LF6‡	116	71	61.2
Average excluding RF3a, RF4:			69.1

Symbols: * only that region of the frame that does not overlap with EC-RF2 is included.

† See text, possibly spliced, but whole frame has been included here.

‡ Only that part of the frame which is in the *EcoRI* C fragment is included.

and also use it to try to detect intervening sequences. This is particularly useful when the intervening sequence is "in frame" with the rest of the coding sequences and contains no termination codons. Below we describe the major features found in the DNA sequence and discuss the possible organization and expression of coding regions.

(3) Organization of reading frames

(a) Region 1-4500

The *in vitro* detected promoter L2 (Farrell *et al.*, 1983) lies within the 3' end of the LF5 open reading frame (Figs 2 and 3). The first methionine codon preceded by a purine at -3 (Kozak, 1981) downstream of this promoter is found at the start of LF6 at position 350, one base beyond the termination codon at the end of LF5. The reading frame extends into the *EcoRI* H fragment and could code for a polypeptide of 717 amino acids (C. Séguin, personal communication). The *EcoRI* H fragment has been shown by sequence analysis to be contiguous with the C fragment. The expression of LF5 would depend on another leftward promoter. The absence of a nearby AATAAA sequence presumably would mean that it would be spliced somewhere at the 3' end. A possible splice donor sequence at position 390 has a seven out of nine base match with the consensus sequence of CAGGTPuAGT .

On the R strand the methionine codon at the beginning of LF5 is overlapped with another methionine codon at the beginning of RF1. This frame remains open until position 2920 overlapping at its 3' end with another open frame RF2, which begins with the methionine codon at 2730. RF2 extends to the termination codon at 4440 and is followed by an AATAAA sequence 43 bases downstream. A possible promoter sequence for RF2 is situated upstream at position 2616 with the sequence

TATTTAAAAA and also the sequence GCCAATT at 2546. Interestingly the 22 base-pairs centred on position 2620 displays twofold symmetry with only two mismatched base-pairs. This potential promoter sequence was not detected by the *in vitro* transcription experiments of Farrell *et al.* (1983), although they did find some low level transcripts emanating from this region. Expression of RF2 could also be effected using the potential splice acceptor sequence two bases upstream of the first methionine codon. No obvious promoter-like sequences can be seen near the beginning of the RF1 and LF5 frames though potential splice acceptor sequences are found at position 2105 for RF1 and at 2244 for LF5. Both of these sequences occur (in the complementary sense) within the coding region of the other reading frame. Thus the possibilities exist for at least the independent expression of RF 2 and possibly of a spliced RF1 + RF 2 frame. In the latter case a good potential splice donor sequence CAG/GTAAGC, which is an eight out of nine match with the donor consensus sequence, exists at position 2256 which is 28 codons downstream of the start of RF1.

(b) *The region 4500-8000*

(i) *RF3 and 3a*

The large open reading frame RF3 extends from the first available methionine codon at 5241 to the termination codon at 7056 which lies five codons past the AATAAA sequence at its 3' end. No *in vitro* promoter was detected upstream of the methionine codon although a potential promoter sequence TATTTAT occurs at 5035. Also possible splice acceptor sequences are found at positions 5059, 5131 and at 5218 before the methionine codon at 5241. There is, however, a promoter in the middle of the RF3 frame with a transcription start at approximately 5962-5972 (Farrell *et al.*, 1983). The first methionine codon is found in frame at position 6021. Thus the potential exists to express the whole frame RF3 or the 3' half RF3a.

(ii) *LF4*

Twenty bases beyond the AATAAA sequence at the 3' end of RF3 is another AATAAA sequence on the L strand at position 7072. Associated with this is an open reading frame LF4 that could code for a protein of 248 amino acids starting at the methionine codon at 7839 and extending to the termination codon at 7095, 28 bases before the AATAAA sequence. The methionine codon is preceded by the sequence:

L strand 5'—CACACCCCACCACATATTTAGG—3'.

The pyrimidine-rich sequence 5' to the AG could serve as a splice acceptor sequence or the underlined sequence could correspond to the "TATA" promoter element. The preceding overlined sequence corresponds to the sequence found in the L2 and R promoters and in a number of other eukaryotic promoters (Farrell *et al.*, 1983). The likely transcription start for this putative promoter would be centred on the 12 base twofold symmetric sequence 5' AGTTTAAAACT 3' at position 7857. The amino acid sequence predicted from the LF4 reading frame contains many potential

glycosylation sites (Neuberger *et al.*, 1972), in fact 12 out of the 14 asparagines in the sequence are followed two residues away with serine or threonine.

(iii) *The B95-8 deletion region*

The B95-8 strain of EBV is missing approximately 13.6 kb of DNA in the *Bam*HI I fragment when compared with other strains (Raab-Traub *et al.*, 1980). Part of the missing DNA has been shown to be homologous to a region of DNA in the *Bam*HI H fragment, which is approximately 90 kb away. We have mapped the deletion point in the B95-8 sequence by also sequencing part of the Raji RI C fragment which does not carry the deletion. So far we only have sequence data for the 5' side of the deleted sequence so we do not yet know whether there are any repeated sequences involved in the deletion event. The data, however, allow us to define accurately the deletion point by comparison of the two sequences. The deletion is between residues 9326-7 of the B95-8 *Eco*RI C fragment. The Raji sequences will be published separately.

The region containing the deletion point between the LF3 and LF4 frames does not contain any open frames longer than 127 codons and we have not yet found any convincing pattern for its expression. An AATAAA sequence occurs on the R strand at 9081 and the largest open reading frame close to this is 64 codons. A possible promoter sequence CATAAAA occurs on the L strand at this point (9094). Other possible promoter sequences occur at 9544 (CATAAAA) and at 9574 (TATAATGA). A repetitive sequence occurs between 8550 and 8940 and the homologous sequences are shown in Table 2. A sequence of 14 consecutive Gs occurs in this region at 8799.

(c) *The region 9470-17,172*

(i) *LF1*

To the right of the deletion point most of the *Eco*RI C fragment sequence is taken up by three large open reading frames LF1-3. These are preceded by the L1 promoter where the transcription start has been mapped to approximately 16,650. The first methionine codon preceded by a purine at -3 occurs at 16,636 at the beginning of the LF1 frame. This frame ends 41 bases before the AATAAA sequence at 14021. However, using the program FRAMESCAN (Staden & McLachlan, 1982) the codon usage pattern changes considerably between 15,550 and 15,490, and to a lesser extent to 15,374, even when using the LF1 codon pattern itself as the input reference sequence. This region contains on the R strand the canonical promoter sequences GGCCAAC (15,464) and TATAAAATAT at (15,518) as noted by Farrell *et al.* (1983) but not detected as an *in vitro* promoter by them. This may indicate that this region is spliced out of the LF1 frame and lends some support to the idea that this region is an *in vivo* rightward promoter that is not detected by the *in vitro* system. There is on the L strand a possible splice donor sequence at 15,601 and possible acceptor sequences at 15,414 and 15,257 that would encompass the potential promoter region. If this region does function as a promoter it would presumably splice to the reading frames downstream at the end of the *Eco*RI C fragment and in the D het fragment in order to

TABLE 2

8550 TTGTTAATCTTTTAGTGGGAAC TAGTGCGGTACCCCTATAGGTCCTACCGAGCTCCTTGCTTGATAATACCCTGTAAACA 8649
*****+***+*****
8672 TTGTTAACCTTTTGGTGSAACTAGTGTAGTGTTGCTGTAAATAAGTGTGCAGGCCACTAGT CACCAGGTGTCAACGGAGGCTACTTTCCTCGAG 8770
*****+***+*****
8814 TCTGTAAACATTTGGTGGCACTGATGTCGTGGTGTGCTGTAATAAAGTGCCTAGCACATCACGTAGGCACCACGAGTGTCACCAG GGCTACTTGCCTCGG 8912

avoid the in phase initiation and termination codons in the 150 bases downstream of the possible transcription initiation point.

(ii) *LF2*

LF2 extends from the methionine codon (14,060) two bases after the LF2 termination codon, past the LF1 AATAAA sequence to the termination codon at 11,015. No candidate for a promoter for LF2 can be predicted so that expression of this frame may be effected by splicing from a leftward promoter such as L1. A possible splice acceptor sequence occurs at 14,137. No AATAAA sequence is found at the 3' end of LF2 so it would presumably be spliced at this end as well. A 22 base-pair sequence with perfect twofold symmetry is found at positions 10,951-10,972 at the end of the LF2 frame.

(iii) *LF3*

This frame lies approximately 600 bases beyond the 3' end of LF2 starting at the methionine at 10,413 and extending to 9475 about 150 bases before the deletion point. No promoter sequences are found but a possible splice acceptor sequence is situated at 10,444. There are two possible initiation codons at the start of the frame at 10,413 and 10,401. No AATAAA sequence is present near the 3' end although this is close to the deletion point. A possible splice donor sequence occurs approximately 50 bases before the end of the frame at 9526. A membrane role is predicted from the amino acid sequence which contains a regular array of hydrophobic regions separated by regions of charged amino acids (see below).

(iv) *RF4*

An open reading frame overlapping LF2 is found on the R strand. No promoter or AATAAA sequences are found near this frame. Several possible donor and acceptor splice sequences are present (13,038, 13,163, 13,190, 13,320 and 13,403) suggesting that perhaps only parts of the overlapping reading frame RF4 might be used.

Although the sequence of the *EcoRI* D het fragment is known (unpublished results) we have not yet overlapped the C and D het sequences to exclude the possibility of a small undetected *EcoRI* fragment between them. Therefore we have not correlated the D het reading frames with those occurring between the L1 promoter and the 3' end of the *EcoRI* C fragment.

(4) *AATAAA sequences*

RF2, RF3, LF1 and LF4 all have AATAAA sequences close to their 3' ends suggesting that these sequences play a role in their mRNA maturation and polyadenylation. Several other AATAAA sequences are found in the *EcoRI* fragment; it is difficult to predict whether these are functional because AATAAA sequences are sometimes found in coding and non-coding sequences of mRNAs. Benoist *et al.* (1980) noted that there was a homologous sequence in the 30 bases downstream of the

AATAAA sequence in some mRNAs with the consensus sequence TTTTCACTGC. Except for a seven out of ten match with a sequence near the AATAAA at 10,573 neither those AATAAA sequences associated with the ends of reading frames nor the isolated AATAAA sequences displayed any homology with this consensus sequence. In some cases possible splice acceptor sequences are seen associated with short reading frames near the other AATAAA sequences. Three such sequences occur upstream from the AATAAA (position 922) at 792, 840 and 877 and in another case at 9049 preceding the AATAAA sequence at 9081. Exceptions to the AATAAA sequences are known; several cases of ATTAAA (listed by Ahmed *et al.*, 1982) and one of ATAA (Setzer *et al.*, 1980). No ATTAAA sequences are found in the *EcoRI* C fragment and we have not searched for the sequence ATAA on the grounds that it is functionally very rare and not unique enough, i.e. not long enough to consider all possible occurrences in this sequence.

(5) Coding capacity

The *EcoRI* C fragment represents approximately 10% of the EBV genome and the sequence analysis of this fragment provides the first opportunity to assess the coding potential and arrangement of the genome. From an analysis of the sequence we would predict at least six to eight proteins coded in this region. However, with multiple choice splicing as found in adenovirus this could be a conservative estimate. The open reading frames cover most of the DNA sequence and their arrangement with respect to each other coupled with their similarity of codon usage (enhanced third position GC) lead us to expect that most if not all of the indicated open frames are expressed into protein. The conclusion is further strengthened by the finding of AATAAA sequences near the 3' ends of half of the large open reading frames which could participate in their mRNA maturation and polyadenylation. The lack of these sequences at the ends of the other open frames coupled with the lack of obvious promoter sequences near the 5' end of most of the frames would suggest that RNA splicing is involved in their putative expression. In most cases there are good candidates for splice acceptor and donor sites near their 5' and 3' ends, respectively. However, given the possible redundancy in these junction sequences, these are difficult to predict with any certainty.

There are several examples of coding economy in the sequence, which is perhaps surprising in such a large viral genome. The termination codon for LF1 and the possible initiation codon for LF2 are only two bases apart. Similarly those for LF5 and LF6 are only one base apart. The possible initiation codons for RF1 and LF5 are overlapping on the opposite strands. The AATAAA sequences associated with the frames RF3, 3a and LF4 which face each other on opposite strands are only 20 bases apart. Thus the mRNA ends would be expected to overlap on the opposite strands by about 20 bases. Also the closeness of the AATAAA sequences to the termination codons are characteristic of the compactness of viral genomes.

(6) Transcription

Three promoters were mapped by Farrell *et al.* (1983) in this fragment using the HeLa cell extract described by Manley *et al.* (1980). This *in vitro* system has been

shown to be an accurate reflection of the *in vivo* situation for some but not all promoters. Thus, for example, late viral promoters depending on a viral function might not be expected to be active in this system. This situation has been argued for another herpes virus HSV1 gene (Frink *et al.*, 1981). Thus it is possible that this region contains other promoters not detected *in vitro*. A number of possibilities have been mentioned previously in the text. If a viral product is indeed required to turn on transcription from these promoters then their detection may depend on the development of an EBV *in vitro* system using a productively EBV infected cell extract.

Recently, Hummel & Kieff (1982a) have mapped a large number of early and late B95-8 RNA transcripts by hybridization to *Bam*HI and *Eco*RI restriction enzyme fragments. In the *Eco*RI C fragment there are five *Bam*HI sites so we can attempt to correlate some of the transcripts with the open reading frames. In *Bam*HI T and X (W in notation of Hummel & Kieff) a 5.0 kb early and a 2.7 kb late transcript have been mapped. These do not extend into the *Bam*HI V fragment (see Fig. 2) (V₁ in their nomenclature) so that it is likely that these transcripts come from the LF5 and LF6 reading frames. RF1 and RF2 are almost totally situated in *Bam*HI V (V₁). Four transcripts are mapped to this fragment of sizes 6.4 kb, 5.5 kb, 3.2 kb and 2.5 kb. As this fragment is only approximately 3 kb in length the relationship of the four transcripts to the RF1 and RF2 frames is unclear. Also these transcripts do not seem to hybridize to the adjacent *Bam*HI fragments, so they could represent multiple splicing events from outside this region to the RF1 and RF2 reading frames. The *Bam*HI d (b) fragment hybridizes to a 1.4 kb late transcript and it is likely that this represents the RF3a reading frame transcribed from the R promoter. *Bam*HI I contains two transcripts, the 1.4 kb late transcript assigned to RF3a and a 1.1 kb late transcript. LF4 and LF3 and part of LF2 are in this fragment. LF3 and LF4 are suitable sizes with LF4 being the best candidate with a possible promoter or splice acceptor sequence at the 5' end and a AATAAA sequence at the 3' end. No other transcripts were detected that could correspond to the large LF2 frame. A 3.0 kb RNA has been mapped to the *Bam*HI A fragment of the *Eco*RI C fragment. This could correspond to the LF1 (857 codons) or the LF2 (1015 codons) frames.

Therefore, although some correspondence can be found between the two sets of mapping data quite a few anomalies remain. It should now be possible to establish an accurate transcription map from this region by a combination of probing the fractionated RNAs with the substantial bacteriophage M13 clone bank of over 500 clones (each containing approximately 400 bases of *Eco*RI C DNA) that were used to generate the *Eco*RI C DNA sequence and by sequencing a cDNA library of the viral transcripts.

(7) Early membrane antigens and the B95-8 deletion

B95-8 has two unique features not found in other EBV strains, the 13.6 kb deletion and different proportions of the membrane antigens gp350 and gp220 (which are antigenically related). There is considerable interest in these membrane antigens because antibodies to them can neutralize EBV infections in tissue culture (Hoffman *et al.*, 1980; Thorley-Lawson & Geilinger, 1980; Thorley-Lawson & Poodry, 1982). These glycoproteins or parts of them might therefore be used for immunizing people against

Whether there is a direct genotype/phenotype relationship between the DNA deletion and altered expression of the membrane antigens is uncertain, but the two leftward reading frames (LF3 and LF4) around the deletion point have features consistent with a membrane glycoprotein. Computerized analysis of the hydropathy (Kyte & Doolittle, 1982) shows that LF3 has alternating stretches of hydrophobic amino acids with charged residues in between and is probably a membrane protein. The most likely arrangement for this polypeptide in a membrane is shown in Figure 4. LF4 has a large number of potential glycosylation sites (sequence Asn-X-Thr/Ser). It is possible that some or all of the leftward reading frames could be spliced together to produce the antigenically related glycoproteins of molecular weights 350,000 and 220,000 (gp350 and gp220). The presence of the deletion between LF3 and LF4 could in some way affect the levels of expression of the reading frames in this region.

[illegible]

Figure 4. Putative folding of the LF3 polypeptide through a lipid bilayer.

complete proteins found *in vivo*. Perhaps by cloning sections of the viral DNA into eukaryotic expression vectors, transfecting those into suitable tissue-culture cells and looking for production of the complete viral protein it will be possible to assign proteins unequivocally to the DNA sequence.

Acknowledgments

During this work P.J.F. held a Medical Research Council training fellowship and P. D. was a fellow of the North Atlantic Treaty Organization.

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